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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.					
10/600,816	06/20/2003	Gena S. Whitney	D0251 NP	5150					
23914	7590 08/01/2006	EXAMINER							
LOUIS J. V	VILLE MYERS SQUIBB COMPA	NV	LI, RUI	XIANG					
	EPARTMENT	14.1	ART UNIT	PAPER NUMBER					
POBOX 40	000		1646						
PRINCETO	N, NJ 08543-4000		DATE MAILED: 08/01/2000	6					

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	Applicant(s)				
		Application No.	Applicant(s)				
	Office Action Summary	10/600,816	WHITNEY ET AL.				
	Onice Action Guinnary	Examiner	Art Unit				
	The MAILING DATE of this communication app	Ruixiang Li	1646				
Period fo		lears on the cover sheet with the c	orrespondence address				
WHIC - Exter after - If NO - Failu Any	ORTENED STATUTORY PERIOD FOR REPLY CHEVER IS LONGER, FROM THE MAILING DATE in an any be available under the provisions of 37 CFR 1.13 SIX (6) MONTHS from the mailing date of this communication. It period for reply is specified above, the maximum statutory period were to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing and patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tirr vill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	I. sely filed the mailing date of this communication. D (35 U.S.C. § 133).				
Status							
1)🖂	Responsive to communication(s) filed on 22 M	ay 2006.					
2a) <u></u> ☐	This action is FINAL . 2b)⊠ This	action is non-final.					
3)□	Since this application is in condition for allowar	·					
	closed in accordance with the practice under E	x parte Quayle, 1935 C.D. 11, 45	53 O.G. 213.				
Dispositi	on of Claims						
5)□ 6)⊠ 7)□	Claim(s) 35-44 is/are pending in the application 4a) Of the above claim(s) 37 and 39-44 is/are with Claim(s) is/are allowed. Claim(s) 35,36 and 38 is/are rejected. Claim(s) is/are objected to. Claim(s) are subject to restriction and/or	vithdrawn from consideration.					
Applicati	on Papers						
10)⊠	The specification is objected to by the Examine The drawing(s) filed on 30 June 2003 is/are: a) Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct The oath or declaration is objected to by the Ex	☑ accepted or b)☐ objected to drawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). sected to. See 37 CFR 1.121(d).				
Priority u	ınder 35 U.S.C. § 119						
a) [Acknowledgment is made of a claim for foreign All b) Some * c) None of: 1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the priority application from the International Bureausee the attached detailed Office action for a list of the certified copies of the certified copies of the priorical detailed Office action for a list of the certified copies of the certified copies of the priorical detailed Office action for a list of the certified copies of the certified copies of the priorical detailed Office action for a list of the certified copies of the certified copies of the priorical detailed Office action for a list of the certified copies of the priorical detailed Office action for a list of the certified copies of the certified copies of the priorical detailed Office action for a list of the certified copies of the certified copies of the priorical detailed Office action for a list of the certified copies of the certifi	s have been received. s have been received in Application rity documents have been received (PCT Rule 17.2(a)).	on No ed in this National Stage				
Attachmen	• *		·				
2) Notic	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449 or PTO/SB/08) r No(s)/Mail Date 5 22 06 7 7 1 22 04	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other: <u>Sequence ali</u>	ate atent Application (PTO-152)				

DETAILED ACTION

Election/Restrictions

1. Applicant's election without traverse of Group II, claim 38, drawn to a method of diagnosing the presence of breast tumor comprising measuring RNA that encodes the polypeptide of SEQ ID NO: 3, in the reply filed on 05/22006 is acknowledged. Claims 35 and 36 are treated as linking claims. Claims 37, and 39-44 have been withdrawn from further consideration pursuant to 37 CFR 1.142(b) as being drawn to a nonelected invention, there being no allowable generic or linking claim.

Information Disclosure Statement

2.The information disclosure statements filed on 05/22/2006, 07/01/2005, and 01/22/2004 have been considered by the Examiner and a signed copy of form PTO-1449 is attached to the office action.

Drawings

3. The drawings filed on 06/03/2003 are accepted by the Examiner.

Objection to Title

4. The title of the invention is not descriptive. A new title is required that is clearly indicative of the invention to which the claims are directed.

Claim Rejections—35 USC § 112, 1st paragraph

5. The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

6. Claims 35, 36, and 38 are rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for a method of diagnosing breast cancer in a sample by determining the expression level of RNA encoding the polypeptide of SEQ ID NO: 3 comprising specific hybridizing between said RNA to the complementary sequence of SEQ ID NO: 2 or its coding sequence, does not reasonably provide enablement for the instantly claimed method comprising hybridizing between said RNA to the complementary sequence of a nucleic acid *comprising* a fragment of SEQ ID NO: 2, a nucleotide sequence encoding the amino acid sequence of SEQ ID NO: 3 or a fragment thereof. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to use the invention commensurate in scope with these claims.

The factors that are considered when determining whether a disclosure satisfies enablement requirement include: (i) the quantity of experimentation necessary; (ii) the amount of direction or guidance presented; (iii) the existence of working examples; (iv) the nature of the invention; (v) the state of the prior art; (vi) the relative skill of those in the art; (vii) the predictability or unpredictability of the art; and (viii) the breadth of the claims. *Ex Parte Forman*, 230 USPQ 546 (Bd Pat. App. & Int. 1986); *In re Wands*, 858 F. 2d 731, 8 USPQ 2d 1400 (Fed. Cir. 1988).

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The linking claims 35 and 36 are drawn to a method of diagnosing the presence of a tumor or predisposition to a tumor in a sample comprising the expression level of RNA encoding a polypeptide comprising the sequence of amino acids 2 to 357 of SEQ ID NO: 3 in a normal tissue sample and in a test tissue sample by measuring RNA of said polypeptide; and comparing said expression level of said polypeptide from said test tissue sample with said expression level of said polypeptide from said normal test sample; wherein an elevated expression level of said polypeptide in said test tissue sample relative to the expression level of said polypeptide in said normal sample is indicative of the presence of a tumor or a predisposition to a tumor. Claim 38 limits the tumor to be a breast tumor.

The claims are broad and are drawn to a method of diagnosing breast cancer using a genus of nucleic acids. While providing sufficient guidance and/or working examples on how to determine the expression level of mRNA encoding the polypeptide of SEQ ID NO: 3 in various normal tissues (Fig. 5) and tumor tissues (breast, stomach tumors, and testicular tumors) (see, e.g., Example 11, Fig. 16-18), using quantitative PCR analysis and specific primers and probe (page 213), the specification fails to provide sufficient guidance/direction or working examples on how to diagnose breast cancer by hybridizing mRNA in a breast tumor sample with a genus of nucleic acids, including a complementary sequence of a nucleic acid comprising a fragment of SEQ ID NO: 2, a nucleotide sequence encoding the amino acid sequence of SEQ ID NO: 3 or a fragment thereof. Thus, use of a complementary sequence of these nucleic acids in the measurement of mRNA level by hybridization

may measure an mRNA that is distinct from the present mRNA. The state of the art is such that determining the specificity of hybridization is empirical by nature and the effect of mismatches is unpredictable, as taught by Wallace et al. (Methods Enzymol. 152:432-443, 1987) and Sambrook et al. (Molecular Cloning, A Laboratory Manual, 2nd Edition, 1989, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY, page 11.47).

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The prior art (see, e.g., U.S. Patent No. 6812339; U.S. Patent Application Publication No. 20030113798A1) teaches an isolated nucleic acid molecule that is 100% identical to SEQ ID NO: 2 and encodes a polypeptide that is 100% identical to SEQ ID NO: 3 of the present invention (see attached sequence alignment). Veiby et al. (U.S. Pub. No. US2003/0068636 A1, April 10, 2003; 102(e) date: 06/21/2001) teach a diagnostic method of assessing whether a patient is afflicted with breast cancer comprising determining the expression level of RNA encoding the polypeptide of SEQ ID NO: 2 (see sequence alignment). However, none of the prior art teaches diagnosing breast cancer by hybridizing mRNA in a breast tumor sample with a complementary sequence of a nucleic acid *comprising* a fragment of SEQ ID NO: 2, a nucleotide sequence encoding the amino acid sequence of SEQ ID NO: 3 or a fragment thereof.

While an artisan has a high level of skill in determining expression profile of an mRNA in normal tissues and tumor samples and diagnosing tumors, such as breast cancer, the recited use of a genus of nucleic acids in the claimed methods would

require an artisan to carry out undue experimentation to practice the claimed invention.

Accordingly, in view of the factors discussed above, it would require undue experimentation for one skilled in the art to use the invention commensurate in scope with these claims.

Claim Rejections 35 USC § 112, 2nd paragraph

7. The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter, which the applicant regards as his invention.

8. Claims 35, 36, and 38 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 35 is indefinite because it, in part b), recites "said expression level of said polypeptide". It is clear from part a) of the claim that the expression level of RNA, not the expression level of polypeptide, is determined.

Claims 36 and 38 are rejected as dependent claims from claim 35.

Claim Rejections—35 USC § 102(e)

9. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(e) the invention was described in (1) an application for patent, published under section 122(b), by another filed in the United States before the invention by the applicant for patent or (2) a patent granted on an application for patent by another filed in the United States before the invention by the applicant for patent,

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except that an international application filed under the treaty defined in section 351(a) shall have the effects for purposes of this subsection of an application filed in the United States only if the international application designated the United States and was published under Article 21(2) of such treaty in the English language.

10. Claims 35, 36, and 38 are rejected under 35 U.S.C. 102(e) as being anticipated by Veiby et al. (U.S. Pub. No. US2003/0068636 A1, April 10, 2003; 102(e) date: 06/21/2001).

Veiby et al. teach a nucleic acid marker (SEQ ID NO: 59) for breast cancer (see Table 2) that comprises the coding sequence of SEQ ID NO: 2 of the present invention and encodes a protein (SEQ ID NO: 60) that is 100% identical to the polypeptide of SEQ ID NO: 3 of the present invention (see attached sequence alignment). Veiby et al. teach a diagnostic method of assessing whether a patient is afflicted with breast cancer comprising determining the level of expression of a marker of the invention in a patient sample and the normal level of expression of the marker in a control non-cancerous breast sample. A significantly higher level of expression of the nucleic acid marker in the patient sample as compared to the normal level is an indication that the patient is afflicted with breast cancer ([0020] to [0023]). Veiby et al. further teach that expression of a nucleic acid marker can be assessed by preparing mRNA/cDNA from cells in a patient sample, and by hybridizing the mRNA/cDNA with a reference polynucleotide which is a complement of a marker nucleic acid, or a fragment thereof ([0122]). Thus, the teachings of Veiby et al. meet the limitations of claims 35, 36, and 38.

Conclusion

11. No claims are allowed.

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Advisory Information

Any inquiry concerning this communication or earlier communications from the

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examiner should be directed to Ruixiang Li whose telephone number is (571) 272-0875.

The examiner can normally be reached on Monday through Friday from 8:30 am to 5:00

pm. If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Gary Nickol, can be reached on (571) 272-0835. The fax number for the

organization where this application or proceeding is assigned is (571) 273-8300.

Information regarding the status of an application may be obtained from the

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applications may be obtained from either Private PAIR or Public PAIR. Status

information for unpublished applications is available through Private PAIR only. For

more information about the PAIR system, see http://pair-direct.uspto.gov. Should you

have guestions on access to the Private PAIR system, please contact the Electronic

Business Center (EBC) at the toll-free phone number 866-217-9197.

Ruixiang L Ruixiang Li, Ph.D.

Primary Examiner

July 29, 2006

RUIXIANG LI, PH.D.

PRIMARY EXAMINER

-continued

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<212> TYPE: DNA

<213> ORGANISM: Homo sapiens

<400> SEQUENCE: 59

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115.2	115.2	115	115	115	114.8	114.8	114.4	113.8	113.8	113.2	113.2	112.6	112.6	112.4	112.4	112.4	112.2	112.2	111.6	111.6	111.6
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ALIGNMENTS

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Sequence 512, Application US/09949016

Sequence 512, Application US/09949016

Patent No. 681239

GENERAL INPORMATION:

PAPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF;

FILE REPERENCE: CL001307

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR FILING DATE: 2000-10-20

PRIOR FILING DATE: 2000-10-03

PRIOR FILING DATE: 2000-10-03

PRIOR FILING DATE: 2000-10-03

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FastSEQ for Windows Version 4.0

SEQ ID NO 572

LENGTH: 2456
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; Pred. No. 0;
0; Mismatches
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Best Local Similarity 100.0%;
Matches 2456; Conservative 0
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US-09-949-016-572
US-09-949-016-572
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KLVRGRKPLSLLVILGLAVGFSLVQDVIALEYIVLTMNRTNVNVFSELSAPRRNEDFVLL 180
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      APPLICANT: G1sh, KUIT C.
APPLICANT: G1sh, KUIT C.
APPLICANT: Hevezi, Peter A.
APPLICANT: Mack, David H.
APPLICANT: Mack, David H.
APPLICANT: Matson, Susan R.
APPLICANT: Watson, Susan R.
APPLICANT: Bos Batocechnology, Inc.
APPLICANT: Bos Batocechnology, Inc.
TITLE OF INVENTION: Methods of Screening for Modulators of Cancer
TITLE OF INVENTION: Methods of Screening for Modulators of Cancer
TITLE OF INVENTION: Methods of Screening for Modulators of Cancer
TITLE OF INVENTION: Methods of Screening for Modulators of Cancer
TITLE OF INVENTION: Methods of Screening for Modulators of Cancer
TITLE OF INVENTION: Methods of Screening for Modulators of Cancer
TITLE OF INVENTION: Methods of Screening for Modulators
CURRENT APPLICATION NUMBER: US 60/350,666
PRIOR FILING DATE: 2001-11-15
PRIOR PELING DATE: 2001-11-21
PRIOR PELING DATE: 2002-01-08
PRIOR PELING DATE: 2002-02-08
PRIOR PELING DATE: 2002-02-08
PRIOR PELING DATE: 2002-02-08
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                                                                                                       100.0%; Score 1865; DB 4;
100.0%; Pred. No. 2.7e-171;
tive 0; Mismatches 0;
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Publication No. US20030232350A1
GENERAL INFORMATION:
APPLICANT: Afar, Daniel
APPLICANT: Aziz, Natasha
APPLICANT: Ginsberg, Wendy M.
APPLICANT: Gish, Kurt C.
                                                                                                            Query Match 100
Best Local Similarity 100.
Matches 357; Conservative
         TYPE: PRT
; ORGANISM: Homo sapiens
US-10-224-289-4
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                                                                                                                        Sequence 49.4, Application US/10225567A

Sequence 49.4, Application US/10225567A

Publication No. US20030113798A1

GENERAL INFORMATION:
APPLICANT: LifeSpan Biosciences
APPLICANT: Burmer, Glenna C.
APPLICANT: Roush, Christine L.
APPLICANT: Roush, Christine L.
TITLE OF INVENTION: ANTIGENIC PEPTIDES AND ANTIBODIES FOR G PROTEIN-COUPLED RECEPTORS
FILE REFERENCE: 1920-4-4

CURRENT PEPLICATION NUMBER: US/10/225,567A

CURRENT PELLING DATE: 2001-12-19

PRIOR FILING DATE: 2000-12-9

NUMBER OF SEQ ID NOS: 2292

SOFTWARE: PatentIn version 3.1

SEQ ID NO 454

LUMBER OF SEQ ID NO 454

LUMBER OF SEQ ID NO 454

LUMBER OF SEQ ID NO 454
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publication No. US20030207288A1

publication No. US20030207288A1

publication No. US20030207288A1

publication No. US20030207288A1

publicant INFORMATION:

predicant: LEMIN, David A.

TYTLE OF INVENTION: GPCR-LIKE RETINOIC ACID-INDUCED GENE 1 PROTEIN AND

TYTLE OF INVENTION: NUCLEIC ACID

TYTLE OF INVENTION: NUCLEIC ACID

TYTLE OF INVENTION: NUCLEIC ACID

CURRENT APPLICATION NUMBER: US/10/224,289

CURRENT FILING DATE: 2001-08-20

PRIOR FILING DATE: 2001-08-20

NUMBER OF SEQ ID NOS: 20

SOFTWARE: Patentin Ver. 2.1

SEQ ID NO 4
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; Pred. No. 2.7e-171;
0; Mismatches 0;
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Best Local Similarity 100.0%;
Matches 357; Conservative 0
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                TYPE: PRT
GRGANISM: Homo sapiens
US-10-225-567A-454
                                                                                                                      -10-225-567A-454
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Published Applications NA Main:*

1: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US07_PUBCOMB.seq:*

2: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US09_PUBCOMB.seq:*

3: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US09_PUBCOMB.seq:*

4: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US09B_PUBCOMB.seq:*

5: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US10B_PUBCOMB.seq:*

6: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US10B_PUBCOMB.seq:*

7: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US10B_PUBCOMB.seq:*

8: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US10B_PUBCOMB.seq:*

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16: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US11B_PUBCOMB.seq:*

16: /EMC_Celerra_SIDS3/ptodata/2/pubpna/US11B_PUBCOMB.seq:*
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Copyright (c) 1993 - 2006 Biocceleration Ltd.
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Sequence 453, App Sequence 63, Appl Sequence 64, Appl Sequence 619, Appl Sequence 2, Appli Sequence 10, Appl Sequence 10424, A Sequence 11, Appl Sequence 3, Appli Sequence 405, App Sequence 9, Appli Sequence 2, Appl Sequence 59, Appl Sequence 40, Sequence 32, Sequence 18, A Description US-10-600-816-18 US-10-198-846-10424 US-10-75-920-11 US-10-224-289-3 US-10-224-289-3 US-10-775-920-9 US-10-775-920-9 US-10-10-775-920-9 US-10-10-6-847-59 US-10-225-567A-453 US-10-269-909-63 US-10-269-909-64 US-10-295-027-619 US-10-600-816-2 US-10-775-920-10 US-10-938-061-40 US-11-169-041-32 Query Match Length DB 2446 100.0 100.0 100.0 100.0 24448 2439.8 2302 2302 2302 2302 2302 2308 Score

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Sequence 59, Appl	Sequence 13, Appl	12,	Sequence 834, App	5, 7	43	64,	64,	2418,	Sequence 6690, Ap	49,	Sequence 249, App	Sequence 210, App	210,	134,	196	23,	Seguence 19238, A	Sequence 2510, Ap	207	Sequence 1, Appli	127	Sequence 2222, Ap		Sequence 1937, Ap	Sequence 1808, Ap	Sequence 396, App	2451,
15 US-11-080-991-59	9 US-10-775-920-13	9 US-10-775-920-12	7 US-10-264-049-834	7 US-10-224-289-5	9 US-10-935-190-43	10 US-10-936-626-64	_	13 US-11-060-756-2418	13 US-11-060-756-6690	3 US-09-866-050A-249	6 US-10-152-661-249	3 US-09-978-360A-210	5 US-09-978-360A-210	9 US-10-712-615-134	10 US-10-505-486-196	7 US-10-313-542-223	3 US-09-864-761-19238	3 US-09-864-761-2510	6 US-10-106-698-2079	7 US-10-224-289-1	3 US-09-969-034-1270	3 US-09-969-034-2222	6 US-10-066-543-1811	6 US-10-066-543-1937	3 US-09-969-034-1808	6 US-10-052-283-396	3 US-09-998-598-2451
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2286	2274.8	2268.2	2260.4	1601.8	1601.8	1441.4	1441.4	1400	1400	1126	1126	1082.6	1082.6	1011	1067.8	1031.6	932.6	787.6	702.4	661.2	602	540.4	538.4	497	496.6	477.2	467
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Sequence 453, Application US/10225567A

Sequence 453, Application US/10225567A

Sequence 453, Application No. US20030113798A1

Sequence 453, Application No. US2004012-19

FILE REFERENCE: 19204-4

CURRENT APPLICATION NUMBER: US/10/225,567A

CURRENT APPLICATION NUMBER: 60/257,144

FRIOR FILING DATE: 2000-12-19

FRIOR FILING DATE: 2000-12-19

NUMBER OF SEQ ID NOS: 2292

SOFTWARE: Patentin version 3.1

SEQ ID NO 453

LENGTH: 2456
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k; Pred. No. 0;
0; Mismatches
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Best Local Similarity 100.0%;
Matches 2456; Conservative 0;
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; ORGANISM: Homo sapiens
US-10-225-567A-453
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	bb 601 TGTCAGTCTGACCTCGTCGGGGGGGGGAGGAAGCCCCTTTCCCTGTTGGTGATTCTGGG 6 w 661 TCTGGCCGTGGGCTTCAGCCTAGTCCAGGATGTTATCGCTATTGAATATATTGTCCTGAC 7 bb 661 TCTGGCCGTGGGCTTCAGCCTAGTCCAGGATGTTATCGCTATTGAATATTGTCCTGAC 7 cy 721 CATGAATAGGACCAATGTCTTTTTTTTGAGTTCCTCGTCGCAATGAAGA 7 ch 721 CATGAATAAGAACCAATGTCTTTTTTTTTCGCTTCGTCGTCGCAATGAAGA 7 ch 721 CATGAATAAGAACCAATGTCTTTTTTTTTTTCGCTTCGTCGTCGCAATGAAGA 7 ch 721 CATGAATAAGAACCAATGTCTTTTTTTTTTTTTTTTTTT	781 CTTGCCCCCCCCCCCCCCCCCCCCCCTCCTCGAGGCCCCCCCC	841 CTCACCTTCTGTGGTTCCTTCACGGCTGGAGACATGGGGCCCACATCTACCTCAC 	Oy 901 GATGCTCCATTGCCATCTGGGTGGCTTGCATCTGCTTGCT		1021 CCTGTTGGCTTATGTTAGTCCCGAGTTTTGGCTGCTCACAAAGCAACGAAACCCCATGGA	OY 1081 TTATCCTGTTGAGGATGCTTTCTGTAAACCTCGACTGAGGAGGGGTATGGTGTGGA 1140	Qy 1141 GAACAGGCCTACTCTCAAGAAAACACTCAAGGTTTTGAAGAGACAGGGACACGCT 1200	OY 1201 CTATGCCCCTATTCCACATTTTCAGCTQCAGAACCAGCCTCCCCAAAAGGAATTCTC 1260 1201 CTATGCCCCTATTCCACACATTTTCAGCTGCAGAACCAGCTCCCCAAAAGGAATTCTC 1260	OY 1261 CATCCACGGGCCCACGCTTCGCCGAGCCCTTACAAAGACTATGAAGTAAAGAAAG	OY 1321 CAGCTAACTCTGAAGAGTGGGACAAATGCAGCGGGGGGGG	OY 1381 TCAAAGGGATGTGGGGAAATCTTGAGTCTTCTGAGAAAACTGRACAAGACACTACGGGA 1440 1381 TCAAAGGGATGTGGGCGAAATCTTGAGTCTTCTGAGAAAACTGTAAAGACACTACGGGA 1440	Oy 1441 ACAGITICCTCCTCCTCCTAACCACAAITCTTCCATGCTGGGCTGATGTGGGCT 1500	Oy 1501 AGRAAGACTCCAGTTCTTAGAGGCGCTGTAGTATTTTTTTTTT	OY 1561 ATACTICITITAAGIGGAGICICAAGCAACICAAGITIAGACCCITACICITITIGET 1	Ov 1621 GTTTTTGAAACAGGATCTTGCTCTGTCAGGCTTGAGTGCAGTGGAGTGCAGATCACAG
2401 GCAATAAAGATGTGGCCACTCTTTCATGGTGGTGGCACCAAAAAAAA	FULT 2 +10-269-909-63 +10-269-909-63 +10-269-909-63 +10-264-909-909 +10-269-909-909-909-909-909-909-909-909-909-9	APPLICANT: IACOBUZIO-DONAHUE, CHRISTINE APPLICANT: MAITRA, ANIRBAN TITAE OF INVENTION: PANCREATIC CANCER DIAGNOSIS AND THERAPIES FILE, REFERENCE: 58303 (71699)	CURRANT APPLICATION NUMBER: US/10/269,909 CURRANT PILING DATE: 2003-10-11 PRIOR APPLICATION NUMBER: 60/328,609 PRIOR ALING DATE: 2001-10-11 PRIOR APPLICATION NUMBER: 60/332,754	PRIOR FYLING DATE: 2001-11-19 NUMBER OR SEQ ID NOS: 87 SOFTWARE:\Patentin Ver. 2.1 EQ ID NO 63 LENGTH: 24,56	TYPE: DNA ORGANISM: Homo sapiens 10-269-909-63	Ouery Match 100.0%; Score 2456; DB 7; Length 2456; Best Local Similarity 100.0%; Pred. No. 0; Matches 2456; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	GGAACTGGAATAGGCGT	TCCTTGTCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCC	ACAACTGCTCAAGTGCGAGGGGGGGATTCCTTCCCTCGGGGGGGG	CTCGCCTGCTCCCCTCTTGCGCGCGGAAGCCACCACCAAGTTCACGGCCAACGCTTGGC [ACTAGGGTCCAGAATGGCTACAACAGTCCCTGATGGTTGCCGCAATGGCCTGAAATCCAA 30 ACTAGGGTTCAAATGGCTACAACAGTCCCTGATGGTTGCCGCAATGGCCTGAAATCCAA 30 ACTAGGGTTCAAAATGGCTACAAAACAAGTACAAATGCAAATGCAAATGAAATCAAAAAAAA	GTACTACAGACTITICGATANGGCTGAAGCTTGGGGCATCGTCTGAAAACGGTGGCCAC	AGCCGGGGTTGTGACCTCGGTGGCTTCATCCTCTCCCGGTCCTCTCCCAC	Treatecteacteteceaatectegeaggicaleaggi	GCGCCATCTTTGGCCTCACCTCGCCTCATCATCGGACTGGGACGGGGGGGG	481 GGGCATCTTTGGCCTCACCTTCGCCTTCGGACTGGGACGGGGCACAGGGCCCAC 540

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Sequence 6443, Ap Sequence 326, App Sequence 326, App Sequence 123, App Sequence 123, App Sequence 54, Appl Sequence 54, Appl Sequence 57, Appl Sequence 6, Appli Sequence 2, Appli Sequence 2, Appli Sequence 349, App Sequence 2, Appli Sequence 2, Appli
                                                                                     8, 2006, 10:32:39; Search time 50 Seconds (without alignments) 624.969 Million cell updates/sec
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/EMC_Celerra_SIDS3/ptodata/2/iaa/7_COMB.pep:*
/EMC_Celerra_SIDS3/ptodata/2/iaa/HCOMB.pep:*
/EMC_Celerra_SIDS3/ptodata/2/iaa/PCTUS_COMB.pep:*
/EMC_Celerra_SIDS3/ptodata/2/iaa/RE_COMB.pep:*
/EMC_Celerra_SIDS3/ptodata/2/iaa/RE_COMB.pep:*
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Biocceleration Ltd.
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US-09-312-283C-326

US-09-312-283C-123

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US-09-964-956-53

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US-09-964-956-34

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US-09-123-756-6

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12 12 12 13 14 15 15 16 16 17 18 18 18 18 18 18 18 18 18 18 18 18 18	TTS METHODS	imilarity 100.0%; Score 1865; DB 2; Length 357; Conservative 0; Mismatches 0; Indels 0; Gaps MattvPDGCRNGLKSKYYRLCDKAEAWGIVLETVATAGVTSVAEMLTLPILVCKVQDSN MATTVPDGCRNGLKSKYYRLCDKAEAWGIVLETVATAGVTSVAEMLTLPILVCKVQDSN RATTVPDGCRNGLKSKYYRLCDKAEAWGIVLETVATAGVTSVAEMLTLPILVCKVQDSN RRWLPTQPLFLLGVLGFGTLFAFIIGLGGSTGPTRFPLFGILFSICFSCLLAHAVSLT RRWLPTQFLFLLGVLGFGLTFAFIIGLDGSTGPTRFPLFGILFSICFSCLLAHAVSLT RRWLPTQFLFLLGVLGFGLFAFIIGLDGSTGPTRFPLFGILFSICFSCLLAHAVSLT KLVRGKRFPLSLLVIIGLAVGFSLVQDVIAIEYIVLTWARTRVNVVFSELSAPRRNEDFVLL KLVRGKKFLSLLVIIGLAVGFSLVQDVIAIEYIVLTWARTRVNVVFSELSAPRRNEDFVLL
US-09-619-353-8 US-08-485-588-5 US-08-484-565-5 US-08-481-751-5 US-08-353-784-5 US-08-353-784-5 US-08-484-719B-5 US-08-484-719B-5 US-08-619-353-7 US-09-619-353-7 US-10-125-772-2 US-10-125-772-2 US-10-125-778-2 US-10-125-778-2 US-10-125-778-2 US-10-125-778-2 US-10-125-778-6 US-10-125-778-6 US-10-125-778-6 US-10-125-778-6 US-10-125-778-6 US-10-125-778-6 US-10-125-778-6	υ, μ	e 1865; smatches sawGrULETY ILIGLOGSTC ILIGL
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	13 Applicat Applicat ATION: TTON: WITON: W WITON: W WITON: W TION WUM S: CLO013 S: CLO013 S: CLO013 CITON WUM ATE: 200 TION WUM ATE: 200 T	arity onserv VPDGCI VPDGCI UPTQFI UPTQFI GRKPL/ GRKPL/
	1016-6443 No. 6443, Application No. 612339 INFORMATION: NO. 61234 INFORMATION: NO. FART: VENTER, J. Cra OF INVENTION: WITH REPERRING: CLO01307 IT APPLICATION NUMBER: TFILING DATE: 2000-1 APPLICATION NUMBER: FILING DATE: 2000-1	Simile 7; CC CC MATTY MATTY REKONI REKONI RELONI KLUNEC
116.5 116 116 116 116 116.5 116.5 115.5 115.5 113.5 11	16-66 6443 N. 6683 N. 6683 N. C. B. C.	tch 357; 357; 1 M 1 M 61 R 61 R 121 K
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254 WVFLLAY
                                                                              TYPE: PRT
ORGANISM: Human
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ORGANISM: Mouse
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US-09-312-283C-326
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WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FIDE REPRENCE: CLOOL307
CURRENT APPLICATION NUMBER: US/09/949,016
PRIOR PILING DATE: 2000-04-14
PRIOR PILING DATE: 2000-10-20
PRIOR PILING DATE: 2000-10-20
PRIOR PILING DATE: 2000-10-3
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PRIOR PILING DATE: 2000-0-03
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PRIOR PILING DATE: 2000-0-09
PRIOR PILING DATE: 2000-0-09
SOFTWARE: PASSECT FOR WINDOWS: 207012
SOFTWARE: PASSECT FOR WINDOWS VERSION 4.0
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US-09-188-330-326
Sequence 326, Application US/0918893AA
Perent No. 6150502
GENERAL INFORMATION:
APPLICANT: Watson, James D.
APPLICANT: Sleeman, Matthew
APPLICANT: Onrust, Rene
APPLICANT: Onrust, Rene
TITLE OF INVENTION: Compositions Isolated Fro;
TITLE OF INVENTION: and Methods For Their Vse
FILE REFERENCE: 11000.1011c1
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CURRENT APPLICATION NUMBER: US/09/188,930A
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99.1%; Pred. No. 3.3e-185;
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CURRENT APPLICATION NUMBER: US/09/312,283C CURRENT FILING DATE: 1999-05-14
NUMBER OF SEQ ID NOS: 425
SEQ ID NO 326
                                                                                                                                                                                                                          2; Mismatches
CURRENT FILING DATE: 1998-11-09
NUMBER OF SEQ ID NOS: 348
SOFTWARE: FastSEQ for Windows Version 3.0
SEQ ID NO 326
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Patent No. 6573095
GENERAL INPORMATION:
APPLICANT: Watson, James D.
APPLICANT: Strachan, Lorna
APPLICANT: Sleeman, Matthew
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341; Conservative
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DDTILSSALAANGWVFLLAYVSPEFWLLTKQRNPMDYPVEDAFCKPQLVKKSYGVENRAY 300
                                                                                                                          The invention comprises a method for assessing whether a patient is afflicted with breast cancer or ovarian cancer. The method involves the use of specific DNA markers. The method of the invention is useful in the detection and treatment of ovarian and breast cancer. Amino acid sequences ABJ37025 - ABJ37080 represent human breast/ovarian cancer-
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Best Local Similarity 100.0%; Pred. No. 2.7e-203;
Matches 357; Conservative 0; Mismatches 0; Indels 0;
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27-JUN-2001; 2001US-0301351P.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              The present invention describes an isolated nucleic acid molecule, which comprises the sequence of any of the genes that are up-regulated or down-egulated in specific cancers (e.g. about 1031 genes up-regulated in acute 1ymphocytic leukemia). ACC72641 to ACC72660 represent cancer related gene nucleotide sequences which encode the proteins given in ABRSB321 to ABRSB709. Also described: (1) determining the presence or absence of a pathological cell in aptient; (2) an expression vector comprising a nucleic acid molecule described above; (3) a host cell comprising the vector; (4) an itsolated polypeptide, which is encoded by the nucleic acid; (5) an antibody that specifically binds the polypeptide of (4); (6) specifically targethy a compound to a pathological cell in a patient by administering to the patient the antibody above; and (7) a patient by administering to the patient the antibody above; and (7) a patient by administering to the patient the antibody above; and (7) a patient by administering to the patient the antibody above; and (7) a patient by administering to the patient the antibody above; and (7) a patient by administering to the patient the nucleic acid is useful for therapeutic targets; In particular, the nucleic acid is useful for bancers, prostate, skin and uterus), wounds, ischaemia, heart disease, atherosclerosis and endometriosis. The nucleic acid is also useful in dry administering these are defined acid in the patient diseases.
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Best Local Similarity 100.0%; Pred. No. 2.7e-203;
Matches 357; Conservative 0; Mismatches 0; Indels 0
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Human cancer related protein SEQ ID NO:226
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2001US-0350666P.
2002US-0355145P.
2002US-0355257P.
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20-SEP 2001; 2
13-NOV-2001; 2
08-FEB-2002; 2
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The present sequence is that of human G-protein coupled receptor-like retinoic acid induced gene 1 (GPCR-like RAIG1) protein. This is the human homologue of murine GPCR-like RAIG1 (see ABR4648). The murine GPCR-like RAIG1 gene was shown to be differentially regulated during fasting-feeding cycles, with moderate induction early in fasting, down-regulation with extended fasting and 4-fold up-regulation with feeding in recovery from fasting. The differentially expressed gene, its mRNA, and the concoded protein, can each be manipulated to detect and treat metabolic disorders associated with up- or down-regulation of GPCR-like RAIG1
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hypertension; hypotension; renal disorder; rheumatoid arthritis; trauma;
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1) polypeptide and gene, useful for diagnosing or treating disorders, e.g. obesity, anorexia, cachexia or diabetes.
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0; Mismatches 0;
                                                                                                                     Disclosure; Page 19-20; 150pp; English.
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